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Submitted for form 1449/PTO				Complete if Known	
				Application Number	10/689,982
INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Filing Date	October 21, 2003
				First Named Inventor	LaColla <i>et al.</i>
				Group Art Unit	1614
				Examiner	Unassigned
Sheet	1	of	3	Attorney Docket Number	06171.105003 IDX 1003 CON

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U.S. PATENT DOCUMENTS						
Examiner Initials *	Cite No. ¹	U.S. Patent Document Number Kind Code ^{2,3} (if known)		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pgs, Clns, Lns, Where Relevant Passages/Relevant Figs Appear
VB	AA	3,956,302	A	Hunter <i>et al.</i>	05-11-1976	
VB	AB	5,747,500	A	Son <i>et al.</i>	05-05-1998	
VB	AC	5,914,402	A	Choi <i>et al.</i>	06-22-1999	
VB	AD	5,998,411	A	Vig <i>et al.</i>	12-07-1999	
VB	AE	6,117,904	A	Murphy <i>et al.</i>	09-12-2000	
VB	AF	6,136,335	A	Uckun <i>et al.</i>	10-24-2000	
VB	AG	6,177,437	B1	Wright	06-23-2001	
VB	AH	6,376,504	B1	Uckun <i>et al.</i>	04-23-2002	
VB	AI	2002/0193415	A1	LaColla <i>et al.</i>	12-19-2002	
VB	AJ	6,545,007	B1	Sommadossi <i>et al.</i>	04-08-2003	
VB	AK	6,635,636	B1	LaColla <i>et al.</i>	10-21-2002	
VB	AL	2003/0225114	A1	Sommadossi <i>et al.</i>	12-04-2003	

FOREIGN PATENT DOCUMENTS								
Examiner Initials *	Cite No. ¹	Foreign Patent Document Office ² Number Kind Code ³ (if known)			Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T 6
W	AM	EP	0,469,685	A1	Shell Int. Res. Maatsch., (Carter <i>et al.</i> , <i>Chem. Abstr.</i> , 116:194341)	02-05-1992		
W	AN	WO	91/18887	A1	Smith-Kline Beecham (Ife <i>et al.</i> , <i>Chem Abstr.</i> , 116:128961)	12-12-1991		
W	AO	WO	92/16201	A1	E.B. Michaels Res. Assoc. Inc.	09-12-2000		
W	AP	WO	95/18109 (PCT/KR94/00178)	A1	Korea Res. Inst. of Chem. Techn.	07-06-1995		
W	AQ	WO	97/43266 (PCT/KR97/00084)	A1	Korea Res. Inst. of Chem. Techn.	11-20-1997		
W	AR	WO	97/44342 (PCT/KR97/00074)	A1	Korea Res. Inst. of Chem. Techn.	11-27-1997		
W	AS	WO	00/03998	A1	Novirio Pharmaceuticals (Idenix)	01-27-2000		

Examiner Signature	V. Balasubramanian	Date Considered	9/4/06
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Sheet	2	of	3	Attorney Docket Number	06171.105003 IDX 1003 CON

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OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS				
Examiner Initials *	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T 6	
W	BA	AROYAN, A.A., <i>et al.</i> , "Pyrimidine derivatives. Substituted 6-(4'-alkoxybenzyl)pyrimidines," <i>Arm. Khim. Zh.</i> , 24(2):161-166 (1971), provided as <i>Chem Abstr.</i> , 75:49022.		
M	BB	ARTICO, M., "Non-Nucleoside Anti-HIV-1 Reverse Transcriptase Inhibitors (NNRTIs): A Chemical Survey From Lead Compounds to Selected Drugs for Clinical Trials," <i>Il Farmaco</i> , 51:305-331 (1996).		
M	BC	ARTICO, M., <i>et al.</i> , "3,4-Dihydro-2-alkoxy-6-benzyl-4-oxypyrimidines (DABO's): A new class of specific inhibitors of human immunodeficiency virus Type 1," <i>Antiviral Chem. Chemother.</i> , 4(6):361-368 (1993).		
M	BD	BABA, M., <i>et al.</i> , "Preclinical evaluation of MKC-442, a highly potent and specific inhibitor of human immunodeficiency virus type 1 <i>in vitro</i> ," <i>Antimicrobial Agents & Chemother.</i> , 38(4):688-692 (April 1994).		
M	BE	BALZARINI, J. <i>et al.</i> , "Human immunodeficiency virus type 1 drug-resistance patterns with different 1-[(2-hydroxyethoxy)methyl]-6-(phenylthio)thymine derivatives," <i>Molecular Pharmacology</i> , 44(4):694-701 (October 1993).		
M	BF	BALZARINI, J. <i>et al.</i> , "Marked inhibitory activity of non-nucleoside reverse transcriptase inhibitors against human immunodeficiency virus type 1 combined with (-)2',3'-dideoxy-3'-thiacytidine," <i>Molecular Pharmacology</i> , 49:882-890(1996).		
M	BG	BOTTA, M., <i>et al.</i> , "Synthesis, antimicrobial and antiviral activities of isotrimethoprim and some related derivatives," <i>Eur. J. Med. Chem.</i> , 27:251-257 (1992).		
M	BH	BROWN, T., <i>et al.</i> , "Isocytosine H2-receptor histamine antagonists. I. Oxmetidine and related compounds," <i>Eur. J. Med. Chem.</i> , 23(1):53-62 (1988), provided as <i>Chem Abstr.</i> , 109:210995.		
M	BI	COSTI, R., <i>et al.</i> , "Structure-activity relationship studies on potential non-nucleoside DABO-like inhibitors of HIV-1 reverse transcriptase," <i>Antiviral Chem. Chemother.</i> , 11(2):117-133 (2000).		
M	BJ	FENNER, H. <i>et al.</i> , "Pyrimido(5,4-B)quinolines," <i>Arch. Pharm.</i> , 311(2):115-125 (1978) (Abstract only; <i>Chem Abstr.</i> , 88(21):152555q).		
M	BK	LIU, X.Y., <i>et al.</i> , "Synthesis and interferon-inducing activity studies on the antiviral compounds of 2,5,6-trisubstituted-4(3H)-pyrimidine derivatives," <i>Yaoxue Xuebao</i> , 29(2):153-157 (1994), shown as <i>Chem. Abstr.</i> , 121:108682.		
M	BL	MAI, A., <i>et al.</i> , "5-Alkyl-2-alkylthio-6-(2,6-dihalophenylmethyl)-3,4-dihydropyrimidin-4(3H)-ones," <i>J. Med. Chem.</i> , 42(4):619-627 (February 25, 1999).		
M	BM	MAI, A., <i>et al.</i> , "Dihydro(alkylthio)(naphthylmethyl)oxypyrimidines: novel non-nucleoside reverse transcriptase inhibitors of the S-DABO series," <i>J. Med. Chem.</i> , 40(10):1447-1454 (May 9, 1997).		
M	BN	MAI, A., <i>et al.</i> , "Synthesis and anti-HIV-1 activity of thioanalogues of dihydroalkoxybenzyloxypyrimidines," <i>J. Med. Chem.</i> , 38(17):3258-3262 (August 18, 1995). XP000578131.		
M	BO	MASSA, S., <i>et al.</i> , "Synthesis and antiviral activity of new 3,4-dihydro-2-alkoxy-6-benzyl-4-oxypyrimidines," <i>Antiviral Chem. Chemother.</i> , 6(1):1-8 (1995). [<i>Chem. Abstr.</i> 122(1):122513c (1995)].		

Examiner Signature	<i>V. Balasubramanian</i>	Date Considered	2/4/02
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OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS			
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M	CA	NIZI <i>et al.</i> , "Solid phase synthesis of 2,6-disubstituted-4(3 <i>H</i>)-pyrimidinones targeting HIV-1 reverse transcriptase," <i>Tetrahedron Letters</i> , 39:3307-3310 (1998).	
M	CB	SBARDELLA, G., <i>et al.</i> , "Does the 2-methylthiomethyl substituent really confer high anti-HIV-1 activity to S-DABOS?" <i>Med. Chem. Res.</i> , 10(1):30-39 (2000).	
M	CC	SBARDELLA, G., <i>et al.</i> , "Structure-activity relationship studies on new DABOS: effect of substitutions at pyrimidine C-5 and C-6 positions on anti-HIV-1 activity," <i>Antiviral Chem. Chemother.</i> , 12(1):37-50 (January 2001).	
M	CD	TANAKA, H., <i>et al.</i> , "Synthesis and antiviral activity of 6-benzyl analogs of 1-[(2-hydroxyethoxy)methyl]-6-(phenylthio)thymine (HEPT) as potent and selective anti-HIV-1 agents," <i>J. Med. Chem.</i> , 38(15):2860-2865 (July 21, 1995).	
M	CE	TRAMONTANO, E., <i>et al.</i> , "Characterization of the anti-HIV-1 activity of 3,4-dihydro-2-alkoxy-6-benzyl-4-oxypyrimidines (DABOs), new non-nucleoside reverse transcriptase inhibitors," <i>Microbiologica</i> , 17(4):269-279 (October 1994).	
M	CF	WAGLE, M.V., <i>et al.</i> , "Tumor inhibitory studies on pyrimidines. 1. 2-Amino-4-hydroxy-5-(β -hydroxyethyl)-6-(alkyl or aryl)pyrimidines," <i>Proc. Indian Acad. Sci., Sect. B</i> , 70(1):9-14 (1969), provided as <i>Chem Abstr.</i> , 71:111298.	

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